



In vivo evaluation of paclitaxel-loaded lipid nanocapsules after intravenous and oral administration on resistant tumor.

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UNLABELLED: Aim & methods: The aim of the present work was to encapsulate paclitaxel (Ptx) in various lipid nanocapsules (LNCs) formulations and then to compare their pharmacokinetics and efficacy on a subcutaneous isograft model in rats.

RESULTS: Three different Ptx formulations were obtained. Drug payloads ranged from 1.32 to 3.62 mg Ptx/g of formulation. After oral administration the area under concentration-time curve was higher ($p < 0.05$) if Ptx was encapsulated, (1,2 Distearoyl-sn-glycero-3-phosphoethanolamine-N-[amino(PEG)] (DSPE-PEG-NH₂)) LNCs displaying the highest area under concentration-time curve ($p < 0.05$).

Efficacy was better than control for standard LNCs after oral administration ($p < 0.05$) and for (DSPE-PEG-NH₂) LNCs after intravenous administration. Despite good absorption, (DSPE-PEG-NH₂) LNCs failed to remain efficient after oral route.

CONCLUSION: This study highlights the importance of efficacy studies paired to pharmacokinetic studies for nanomedicines.

Résumé en anglais

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